### PATENT COOPERATION TREATY

## **PCT**

# INTERNATIONAL PRELIMINARY REPORT ON PATENTABILITY (Chapter I of the Patent Cooperation Treaty)

(PCT Rule 44bis)

Applicant's or agent's file reference 701039-54481	FOR FURTHER ACTION	See item 4 below			
International application No. PCT/US2005/000714	International filing date (day/month/year) 10 January 2005 (10.01.2005)	Priority date (day/month/year) 09 January 2004 (09.01.2004)			
International Patent Classification (8th edition unless older edition indicated) See relevant information in Form PCT/ISA/237					
Applicant CHILDREN'S MEDICAL CENTER COPORATION					

1.	. This international preliminary report on patentability (Chapter I) is issued by the International Bureau on behalf of the International Searching Authority under Rule 44 <i>bis</i> .1(a).				
2.	This REPORT consists of a total of 4 sheets, including this cover sheet.				
	In the attached sheets, any reference to the written opinion of the International Searching Authority should be read as a reference to the international preliminary report on patentability (Chapter I) instead.				
3. This report contains indications relating to the following items:					
	Box No. I	Basis of the report			
	Box No. II	Priority			
	Box No. III	Non-establishment of opinion with regard to novelty, inventive step and industrial applicability			
	Box No. IV	Lack of unity of invention			
	Box No. V	Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement			
	Box No. VI	Certain documents cited			
	Box No. VII	Certain defects in the international application			
	Box No. VIII	Certain observations on the international application			
4. The International Bureau will communicate this report to designated Offices in accordance with Rules 44bis.3(c) and 93bis.1 but not, except where the applicant makes an express request under Article 23(2), before the expiration of 30 months from the priority date (Rule 44bis .2).					
		Date of issuance of this report 10 July 2006 (10.07.2006)			
		10 July 2006 (10.07.2006)			

Authorized officer

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The International Bureau of WIPO 34, chemin des Colombettes 1211 Geneva 20, Switzerland

### PATENT COOPERATION TREATY

From the INTERNAT	IONAL SEARC	HING AUTH	ORITY		REC'D 0 3 JUN 2005
To: DAVID S. RESNICK NIXON PEABODY LLP 100 SUMMER STREET BOSTON, MA 02110-2131		РСТРО РСТ			
		WRITTEN OPINION OF THE INTERNATIONAL SEARCHING AUTHORITY			
					(PCT Rule 43bis.1)
		<del> </del>		Date of mailing (day/month/year)	01 JUN 2005
Applicant'	s or agent's file r	eference		FOR FURTHER ACTION See paragraph 2 below	
701039-54	1481 nal application No	· · · · · · · · · · · · · · · · · · ·	International filing date		
PCT/US05		,,	10 January 2005 (10.01.		
		cation (IPC)	or both national classification		09 January 2004 (09.01.2004)
; C07K 1/0	6/63, 64, 174; 53 00, 14/00, 16/00,		86, 387.1, 387.7, 387.9, 38	8.1 and US Cl.: GO1	N 1/00, 33/48; C07K 1/00, 14/00, 17/00; A61K
Applicant CHILDRE	N'S MEDICAL (	CENTER CO	RPORATION		
1. This o	pinion contains in	ndications rela	ating to the following item	s:	
	Box No. I	Basis of the opinion			
	Box No. II	Priority			
	Box No. III	Non-establi	shment of opinion with reg	gard to novelty, inver	tive step and industrial applicability
	Box No. IV	Lack of uni	ty of invention		
	Box No. V	Reasoned statement under Rule 43bis.1(a)(i) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement			
	Box No. VI	Certain doc	uments cited		
	Box No. VII	Certain defe	ects in the international app	olication	
	Box No. VIII	Certain obs	ervations on the internation	nal application	
2. FUR'	THER ACTIO	N			
If a demand for international preliminary examination is made, this opinion will be considered to be a written opinion of the International Preliminary Examining Authority ("IPEA") except that this does not apply where the applicant chooses an Authority other than this one to be the IPEA and the chosen IPEA has notified the International Bureau under Rule 66.1bis(b) that written opinions of this International Searching Authority will not be so considered.					
If this opinion is, as provided above, considered to be a written opinion of the IPEA, the applicant is invited to submit to the IPEA a written reply together, where appropriate, with amendments, before the expiration of 3 months from the date of mailing of Form PCT/ISA/220 or before the expiration of 22 months from the priority date, whichever expires later.					
For further options, see Form PCT/ISA/220.					
3. For further details, see notes to Form PCT/ISA/220.					
Name and mailing address of the ISA/ US		Authorized officer)			
Mail Stop PCT, Attn: ISA/US Commissioner for Patents		Alana M. Harrisch, Bull Jack on			
P.O. Box 1450 Alexandria, Virginia 22313-1450 Facsimile No. (703) 305-3230				Telephone No. 571-272-1600	

Facsimile No. (703) 305-3230
Form PCT/ISA/237 (cover sheet) (January 2004)

# WRITTEN OPINION OF THE INTERNATIONAL SEARCHING AUTHORITY

International application No.

PCT/US05/00714

Box No. I Basis of this opinion				
1. With regard to the language, this opinion has been established on the basis of the international application in the language in which it was filed, unless otherwise indicated under this item.				
This opinion has been established on the basis of a translation from the original language into the following language, which is the language of a translation furnished for the purposes of international search (under Rules 12.3 and 23.1(b)).				
2. With regard to any nucleotide and/or amino acid sequence disclosed in the international application and necessary to the claimed invention, this opinion has been established on the basis of:				
a. type of material				
a sequence listing				
table(s) related to the sequence listing				
b. format of material				
in written format				
in computer readable form				
c. time of filing/furnishing				
contained in international application as filed.				
filed together with the international application in computer readable form.				
furnished subsequently to this Authority for the purposes of search.				
3. In addition, in the case that more than one version or copy of a sequence listing and/or table relating thereto has been filed or furnished, the required statements that the information in the subsequent or additional copies is identical to that in the application as filed or does not go beyond the application as filed, as appropriate, were furnished.				
4. Additional comments:				
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## WRITTEN OPINION OF THE INTERNATIONAL SEARCHING AUTHORITY

International application No. PCT/US05/00714

Box No. V Reasoned statement under Rule 43 bis.1(a)(i) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement

1. Statement	-	-
Novelty (N)	Claims 17-19	YES
	Claims 1-16	NO
Inventive step (IS)	Claims NONE	YES
	Claims <u>1-19</u>	NO
Industrial applicability (IA)	Claims 1-19	YES
	Claims NONE	NO

#### 2. Citations and explanations:

Claims 1-16 lack novelty under PCT Article 33(2) as being anticipated by Roy et al. (The Journal of Biological Chemistry 279(49): 51323-51330, December 3, 2004). Roy teaches the detection of urinary ADAM 12 in samples from breast cancer patients and controls analyzed by immunoblot, see abstract. ADAM 12 protein levels were higher in urine from breast cancer patients than in control urine. Median levels of ADAM 12 in urine significantly increases with disease progression, see page 51324, column 1, first paragraph; bridging paragraph of pages 51325 and 51326; page 51329, column 2, first two sentences of last paragraph. Western blot analysis was conducted using labeled proteins and chemiluminescence, page 51324, column 2, Western...section.

Claims 17-19 lack an inventive step under PCT Article 33(3) as being obvious over Claims 1-16 lack novelty under PCT Article 33(2) as being anticipated by Roy et al. (The Journal of Biological Chemistry 279(49): 51323-51330, December 3, 2004). The teachings of Roy are presented above. Although Roy et al. does not teach a kit for detecting ADAM 12 in a urine sample comprising a container, at least one antibody and directions for use a scientist would have been inclined to package these components, as well as other reagents that would aid in cancer diagnosis for convenience.